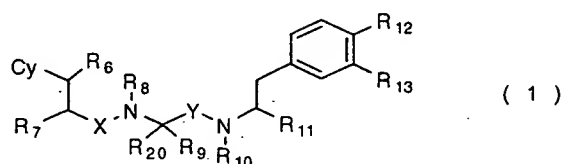


Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

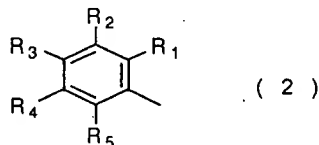
Listing of Claims:

1. (Currently Amended) A compound of Formula (1):



wherein:

Cy is a group of Formula (2):



~~C<sub>3</sub>-cycloalkyl or phenyl,~~

~~R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub> and R<sub>5</sub> are hydrogen, halogen, or hydroxy, amino, trifluoromethyl or nitrile and at least one of R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub> and R<sub>5</sub> is halogen, trifluoromethyl or nitrile;~~

~~R<sub>6</sub> is hydrogen, optionally substituted straight-chained or branched C<sub>1-3</sub> alkyl, amino or hydroxy;~~

~~R<sub>7</sub> is hydrogen, optionally substituted straight-chained or branched C<sub>1-3</sub>alkyl, substituted with one or more hydroxyl groups, or amino optionally substituted with one or~~

~~more of the same or different kinds of straight-chained or  
branched C<sub>1-3</sub> alkyl groups which may be the same or different,  
or hydroxy;~~

R<sub>8</sub> is hydrogen, methyl or ethyl;

R<sub>9</sub> is ~~optionally substituted~~ straight-chained or  
branched C<sub>1-6</sub> alkyl optionally substituted with one or more  
groups which may be the same or different and are selected  
from the group consisting of phenyl, para-hydroxyphenyl, para-  
fluorophenyl, para-chlorophenyl, C<sub>3-7</sub> cycloalkyl, halogen and  
thienyl, ~~optionally substituted straight chained or branched  
C<sub>2-6</sub>alkenyl, optionally substituted straight chained or  
branched C<sub>2-6</sub>alkynyl, C<sub>3-7</sub>cycloalkyl; or optionally substituted  
phenyl;~~

R<sub>20</sub> is ~~hydrogen or straight chained or branched  
C<sub>1-3</sub>alkyl or R<sub>9</sub> and R<sub>20</sub> may together form C<sub>3-7</sub>cycloalkyl;~~

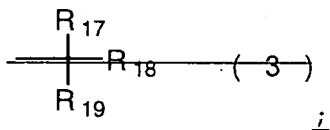
R<sub>10</sub> is hydrogen or methyl or ethyl ~~straight chained  
or branched C<sub>1-3</sub>alkyl;~~

R<sub>11</sub> is ~~hydrogen,~~ straight-chained or branched C<sub>1-3</sub>  
alkyl optionally substituted with one or more groups which may  
be the same or different and are selected from the group  
consisting of ~~amino optionally substituted with one or more of  
the same or different straight chained or branched C<sub>1-3</sub> alkyl,  
3 to 7 membered cyclic amino optionally substituted with  
hydroxyl, amino, carboxyl, carbamoyl or methyl; hydroxyl,~~

~~methoxy, halogen, carbamoyl, methanesulfonyl, ureide,~~  
~~guanidyl, N'-cyano-N''-methylguanidyl, sulfamoylamino,~~  
~~carbamoylmethylamino and methanesulfonylamino, and -CO-~~  
~~N(R<sub>14</sub>)R<sub>15</sub>-carboxyl;~~

~~R<sub>12</sub> is hydroxy or OR<sub>16</sub>;~~

~~R<sub>13</sub> is hydrogen, straight-chained or branched C<sub>1-6</sub>~~  
~~alkyl, straight-chained or branched C<sub>2-6</sub>alkenyl, straight-~~  
~~chained or branched C<sub>2-6</sub>alkynyl or a group of Formula (3):~~



~~R<sub>14</sub> and R<sub>15</sub>, which may be the same or different, are~~  
~~each hydrogen, straight-chained or branched C<sub>1-3</sub> alkyl~~  
~~optionally substituted with straight-chained or branched C<sub>1-3</sub>~~  
~~alkoxy optionally substituted with hydroxyl, amino, carboxyl~~  
~~or carbamoyl, hydroxyl, amino, methylamino, dimethylamino,~~  
~~carbamoyl or methanesulfonyl; optionally substituted straight-~~  
~~chained or branched C<sub>1-4</sub> alkyl, C<sub>3-7</sub>cycloalkyl, straight-chained~~  
~~or branched C<sub>1-4</sub> alkoxy, straight-chained or branched C<sub>1-4</sub>~~  
~~alkylsulfonyl, or pyridyl; a heterocyclic ring;~~

~~R<sub>16</sub> is straight-chained C<sub>1-4</sub> alkyl;~~

~~R<sub>17</sub> is hydrogen or methyl;~~

~~R<sub>18</sub> and R<sub>19</sub> together form cycloalkyl or C<sub>3-7</sub>~~  
~~cycloalkenyl;~~

X is carbonyl or methylene;

Y is carbonyl ~~or~~ methylene;

or a pharmaceutically acceptable salt thereof.

2. (Previously presented) The compound according to claim 1,  
wherein Cy in Formula (1) is a group of Formula (2);  
or a pharmaceutically acceptable salt thereof.

3. (Previously presented) The compound according to claim 1,  
wherein Cy in Formula (1) is a group of Formula (2) in which  
at least one of R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub> and R<sub>5</sub> is halogen and the others  
are hydrogen or hydroxy;  
or a pharmaceutically acceptable salt thereof.

4. (Previously presented) The compound according to claim 1,  
wherein Cy in Formula (1) is a group of Formula (2) in which  
R<sub>3</sub> is halogen or R<sub>2</sub> and R<sub>3</sub> are the same kind of halogen;  
or a pharmaceutically acceptable salt thereof.

5. (Previously presented) The compound according to claim 1,

wherein Cy in Formula (1) is a group of Formula (2) in which R<sub>3</sub> is halogen and R<sub>1</sub>, R<sub>2</sub>, R<sub>4</sub> and R<sub>5</sub> are hydrogen, or R<sub>2</sub> and R<sub>3</sub> are the same kind of halogen and R<sub>1</sub>, R<sub>4</sub> and R<sub>5</sub> are hydrogen; or a pharmaceutically acceptable salt thereof.

Claims 6-13. (Canceled)

14. (Previously presented) The compound according to claim 1, wherein R<sub>7</sub> in Formula (1) is hydrogen or amino optionally substituted with one or more of the same of different kinds of straight-chained or branched C<sub>1-3</sub> alkyl; or a pharmaceutically acceptable salt thereof.

15. (Previously presented) The compound according to claim 1, wherein R<sub>8</sub> in Formula (1) is hydrogen or methyl; or a pharmaceutically acceptable salt thereof.

16. (Previously presented) The compound according to claim 1, wherein R<sub>9</sub> in Formula (1) is methyl, isopropyl, isobutyl, sec-butyl, tert-butyl, 3-pentyl, neopentyl, cyclohexyl, phenyl, benzyl, para-hydroxybenzyl, cyclohexylmethyl or para-fluorobenzyl; or a pharmaceutically acceptable salt thereof.

Claims 17-18. (Cancelled)

19. (Currently Amended) The compound according to claim 1, wherein R<sub>11</sub> in Formula (1) is methyl, hydroxymethyl,

carbamoylmethyl, methanesulfonylmethyl, ureidemethyl,  
sulfamoylaminomethyl, methanesulfonylaminomethyl, ~~earbamoyl~~,  
ethylcarbamoyl, n-propylcarbamoyl, isopropylcarbamoyl,  
~~cyclopropylcarbamoyl~~, tertbutylcarbamoyl, methoxycarbamoyl,  
methylcarbamoyl, methanesulfonylmethylcarbamoyl,  
methoxymethylcarbamoyl,;

or a pharmaceutically acceptable salt thereof.

Claim 20 Cancelled

21. (Currently Amended) The compound according to  
claim 1, wherein  $R_{13}$  in Formula (1) is isopropyl, tert-butyl  
(tBu), or 1,1-dimethylpropyl ~~or 1,1-dimethyl-2-propenyl~~;  
or a ~~hydrate or~~ pharmaceutically acceptable salt thereof.

22. (Currently Amended) The compound according to  
claim 1, wherein in Formula (1) Cy is a group of Formula (2)  
in which at least one of  $R_1$ ,  $R_2$ ,  $R_3$ ,  $R_4$  and  $R_5$  is halogen and  
the others are hydrogen or hydroxy;

~~$R_6$  is hydrogen or methyl;~~

~~$R_7$  is hydrogen or optionally substituted amino optionally  
substituted with one or more of the same or different  
straight chained or branched  $C_{1-3}$  alkyl;~~

$R_8$  is hydrogen or methyl;

R<sub>9</sub> is methyl, isopropyl, isobutyl, sec-butyl, tert-butyl, 3-pentyl, neopentyl, cyclohexyl, phenyl, ~~benzyl, para-hydroxybenzyl, para-fluorobenzyl or cyclohexylmethyl;~~  
~~R<sub>20</sub> is hydrogen;~~  
~~R<sub>10</sub> is hydrogen or methyl;~~  
R<sub>11</sub> is methyl, hydroxymethyl, carbamoylmethyl, methanesulfonylmethyl, ureidemethyl, sulfamoylaminomethyl, methanesulfonylaminomethyl, ~~carbamoyl,~~ methylcarbamoyl, ethylcarbamoyl, n-propylcarbamoyl, isopropylcarbamoyl, ~~tert-cyclopropylcarbamoyl butylcarbamoyl,~~ , methanesulfonylmethylcarbamoyl, methoxymethylcarbamoyl, or methoxycarbamoyl;  
~~R<sub>12</sub> is hydroxy;~~  
R<sub>13</sub> is isopropyl, tert-butyl (tBu), 1,1-dimethylpropyl-or 1,1-dimethyl-2-propenyl;  
or a pharmaceutically acceptable salt thereof.

23. (Previously presented) The compound according to claim 1 which is selected from the group of compounds consisting of Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH<sub>2</sub>, Phe(4-Cl)-N-Me-Val-N-Me-Tyr(3-tBu)-NH<sub>2</sub>, Phe(3,4-F<sub>2</sub>)-N-Me-Val-N-Me-Tyr(3-tBu)-NH<sub>2</sub>, Phe(3-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH<sub>2</sub>, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHOMe, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-tertbutyl-4-hydroxyphenyl)-1-(2-

pyridylcarbamoyl)ethylamide, N-(2-(2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methyl-butyrylamino)-3-(3-tBu-4-hydroxyphenyl)propyl)urea, N-(2-(2-(2-amino-3-(4-fluorophenyl)propanoyl-N-methylamino)-3-methyl)butyrylamino)-3-(3-tertbutyl-4-hydroxyphenyl)propyl)sulfamide, N-[2-(3-tertbutyl-4-hydroxyphenyl)-1-(methanesulfonylaminomethyl)ethyl]-2-[N-(4-fluorophenylalanyloyl)methylamino]-3-methylbutanamide, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-t-butyl-4-hydroxyphenyl)-1-carbamidomethylethylamide, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-t-butyl-4-hydroxyphenyl)-1-methanesulfonylmethylethylamide, 2-(2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methyl-butyrylamino)-3-(3-tBu-4-hydroxyphenyl)propanol, 2-(1-(2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methyl-butyrylamino)-2-(3-tertbutyl-4-hydroxyphenyl)ethyl)-6-methyl-4-pyrimidinone, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-t-butyl-4-hydroxyphenyl)-1-(1,3,4-oxadiazol-2-yl)ethylamide, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-t-butyl-4-hydroxyphenyl)-1-(1,2,4-oxadiazol-5-yl)ethylamide, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-



methylbutyric acid 2-(3-tertbutyl-4-hydroxyphenyl)-1-(thiazol-2-yl)ethylamide, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-t-butyl-4-hydroxyphenyl)-1-(1,3,4-triazol-2-yl)ethylamide, Tyr(2-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH<sub>2</sub>, Tyr(3-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH<sub>2</sub>, Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NH<sub>2</sub>, N-Me-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NH<sub>2</sub>, N-Et-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NH<sub>2</sub>, Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHMe, N-Me-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHMe, N-Et-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHMe, N-Me-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH<sub>2</sub>, N-Et-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH<sub>2</sub>, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHMe, N-Me-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHMe, N-Et-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHMe, Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NH<sub>2</sub>, N-Me-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NH<sub>2</sub>, N-Et-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NH<sub>2</sub>, Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NHMe, N-Me-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NHMe, Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHtBu, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHCH<sub>2</sub>SO<sub>2</sub>CH<sub>3</sub>, Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHEt, N-Me-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHEt, N-Et-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHEt, Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHCH<sub>2</sub>OH, N-Me-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHCH<sub>2</sub>OH, N-Et-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHCH<sub>2</sub>OH, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHEt, N-Me-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHEt, N-Et-Phe(4-F)-N-Me-Val-N-Me-

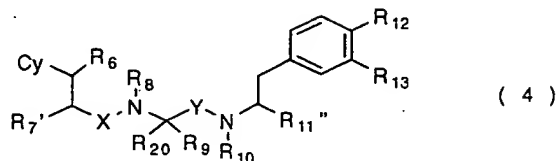
Tyr(3-tBu)-NH<sub>2</sub>Et, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHCH<sub>2</sub>OH,  
N-Me-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHCH<sub>2</sub>OH, N-Et-Phe(4-F)-  
N-Me-Val-N-Me-Tyr(3-tBu)-NHCH<sub>2</sub>OH, Phe(4-F)-N-Me-Val-N-Et-  
Tyr(3-tBu)-NH<sub>2</sub>Et, N-Me-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NH<sub>2</sub>Et,  
N-Et-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NH<sub>2</sub>Et, Phe(4-F)-N-Me-  
Val-N-Et-Tyr(3-tBu)-NHCH<sub>2</sub>OH, N-Me-Phe(4-F)-N-Me-Val-N-Et-  
Tyr(3-tBu)-NHCH<sub>2</sub>OH, N-Et-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-  
NHCH<sub>2</sub>OH, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH<sub>2</sub>Pr, and Phe(4-  
F)-N-Me-Val-Tyr(3-tBu)-NH<sub>2</sub>nPr Phe(4-F)-N-Me-Val-Tyr(3-tBu)-  
NH<sub>2</sub>iPr;  
or a pharmaceutically acceptable salt thereof.

24. (Previously Presented) A pharmaceutical composition containing an effective amount of the compound according to claim 1 as an active ingredient and an inert pharmaceutically acceptable carrier.

25. (Previously Presented) A motilin receptor antagonist composition containing an effective amount of the compound according to claim 1 and an inert pharmaceutically acceptable carrier.

Claims 26-27. (Cancelled)

28. (Currently Amended) A compound of Formula (4):



wherein

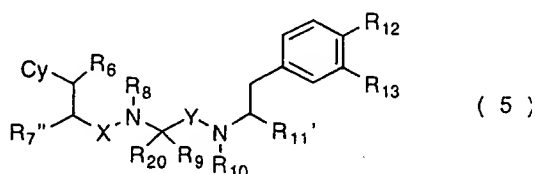
Cy, R<sub>6</sub>, R<sub>8</sub>, R<sub>9</sub>, R<sub>20</sub>, R<sub>10</sub>, R<sub>12</sub>, R<sub>13</sub>, X and Y are as defined in claim 1;

R<sub>7</sub>' is ~~hydrogen, straight-chained or branched C<sub>1-3</sub> alkyl substituted with one or more optionally having at least one-protected hydroxyl groups substituent, or protected amino optionally substituted with having at least one or more substituent of the same or different straight-chained or branched C<sub>1-3</sub> alkyl groups which may be the same or different or protected hydroxyl; and~~

R<sub>11</sub>" is ~~hydrogen, straight-chained or branched C<sub>1-3</sub> alkyl optionally substituted with one or more groups which may be the same or different and are selected from the group consisting of amino optionally substituted with one or more of the same of different straight chained or branched C<sub>1-3</sub> alkyl, 3 to 7 membered cyclic amino optionally substituted with hydroxyl, amino, carboxyl, carbamoyl or methyl, hydroxyl, methoxy, halogen; carbamoyl, methanesulfonyl, ureide, guanidyl, N'-cyano-N''-methylguanidyl, sulfamoylamino, carbamoylmethylamino, and methanesulfonylamino, and -CO-~~

$N(R_{14})R_{15}$ , wherein  $R_{14}$  and  $R_{15}$  are as defined in claim 1,  
~~carboxyl, straight chained or branched  $C_{1-3}$  alkyl having a~~  
~~protected amino;~~  
or a pharmaceutically acceptable salt thereof.

29. (Currently Amended) A compound of Formula (5):



wherein:

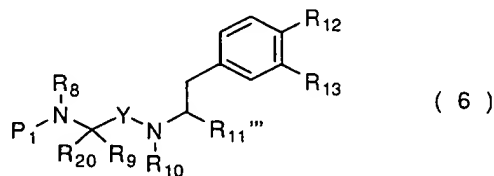
$Cy$ ,  $R_6$ ,  $R_8$ ,  $R_9$ ,  $R_{20}$ ,  $R_{10}$ ,  $R_{12}$ ,  $R_{13}$ ,  $X$  and  $Y$  are as defined in claim 1;

$R_7''$  is ~~hydrogen, straight-chained or branched  $C_{1-3}$  alkyl optionally having at least optionally substituted with one or more optionally protected hydroxyl groups substituent or amino optionally having at least substituted with one or more substituents which are the same or different straight-chained or branched  $C_{1-3}$  alkyl groups which may be the same or different, or optionally protected hydroxy;~~ and

$R_{11}'$  is ~~hydrogen, straight-chained or branched  $C_{1-3}$  alkyl optionally substituted with one or more groups having at least one protected substituent s~~ which may be the same or different and are selected from the group consisting of protected amino ~~optionally substituted with one or more~~

~~straight-chained or branched C<sub>1-3</sub> alkyl, protected 3 to 7-~~  
~~membered cyclic amino optionally substituted with protected~~  
~~hydroxyl, protected amino, protected carboxyl or protected~~  
~~carbamoyl; protected hydroxyl<sub>1</sub>, protected carbamoyl<sub>1</sub>,~~  
~~protected ureide<sub>1</sub>, protected guanidyl<sub>1</sub>, protected N'-cyano-~~  
~~N''-methylguanidyl<sub>1</sub>, protected sulfamoylamino<sub>1</sub>, protected~~  
~~carbamoylmethylamino and protected methanesulfonylamino; and~~  
~~-CO-N(R<sub>14</sub>)R<sub>15</sub> wherein R<sub>14</sub> and R<sub>15</sub> are as those defined in claim 1~~  
~~which are appropriately protected, carboxyl~~  
or a ~~hydrate or~~ pharmaceutically acceptable salt thereof.

30. (Currently Amended) A compound of Formula (6):



wherein:

R<sub>8</sub> is hydrogen, methyl or ethyl~~optionally~~  
~~substituted straight-chained or branched C<sub>1-3</sub> alkyl, optionally~~  
~~substituted amino, or hydroxy;~~

R<sub>9</sub>, is ~~optionally substituted~~ straight-chained or  
branched C<sub>1-6</sub> alkyl optionally substituted with one or more  
groups which may be the same or different and are selected  
from the group consisting of phenyl, para-hydroxyphenyl, para-  
fluorophenyl, para-chlorophenyl, C<sub>3-7</sub> cycloalkyl, halogen and

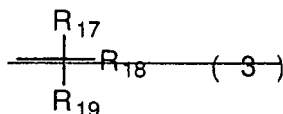
~~thienyl, optionally substituted straight chained or branched  
C<sub>2-6</sub> alkenyl, optionally substituted straight chained or  
branched C<sub>2-6</sub> alkynyl, C<sub>3-7</sub> cycloalkyl or optionally substituted  
phenyl;~~

~~R<sub>20</sub> is hydrogen or methyl or straight chained or  
branched C<sub>1-3</sub> alkyl; or R<sub>9</sub> and R<sub>20</sub> may together form C<sub>3-7</sub>  
cycloalkyl;~~

~~R<sub>10</sub> is hydrogen or methyl or ethyl straight chain or  
branched C<sub>1-3</sub> alkyl;~~

~~R<sub>12</sub> is hydroxy or OR<sub>16</sub>;~~

~~R<sub>13</sub> is hydrogen, straight-chained or branched C<sub>1-6</sub>  
alkyl, straight chained or branched C<sub>2-6</sub> alkenyl, straight-  
chained or branched C<sub>2-6</sub> alkynyl or a group of Formula (3)~~



~~Wherein R<sub>17</sub> is hydrogen or methyl;~~

~~R<sub>18</sub> and R<sub>19</sub> together form cycloalkenyl or C<sub>3-7</sub>  
cycloalkenyl; and~~

~~Y is carbonyl or methylene;~~

~~P<sub>1</sub> is hydrogen or a protecting group of amine; and~~

~~R<sub>11'''</sub> is hydrogen, straight-chained or branched C<sub>1-3</sub>alkyl,  
carboxyl, straight-chained or branched C<sub>1-3</sub>alkyl optionally  
substituted with one or more groups which may be the same or  
different and are selected from the group consisting of amino~~

~~optionally substituted with one or more of the same or different straight chained or branched C<sub>1-3</sub> alkyl, 3 to 7 membered cyclic amino optionally substituted with hydroxyl, amino, carboxyl, carbamoyl or methyl, hydroxyl, methoxy, halogen, carbamoyl, methanesulfonyl, ureide, guanidyl, N'-cyano-N''-methylguanidyl, sulfamoylamino, carbamoylmethylamino and methanesulfonylamino; carboxyl, straight-chained or branched C<sub>1-3</sub> alkyl having protected amino or an optionally substituted heterocyclic ring, or and -CO-N(R<sub>14</sub>)R<sub>15</sub> wherein R<sub>14</sub> and R<sub>15</sub>, which may be the same or different, are hydrogen, optionally substituted straight-chained or branched C<sub>1-4</sub> alkyl optionally substituted with hydroxy, C<sub>3-7</sub> cycloalkyl, straight-chained or branched C<sub>1-4</sub> alkoxy, straight-chained or branched C<sub>1-4</sub>alkylsulfonyl, or pyridyl a heterocyclic ring, carboxyl, straight chained or branched C<sub>1-3</sub>alkyl having protected amino or an optionally substituted heterocyclic ring;~~  
or a pharmaceutically acceptable salt thereof.

Claims 31-34. (Canceled)

35. (Previously Presented) The compound according to claim 1, wherein the substitution of the optionally substituted straight-chained or branched C<sub>1-3</sub> alkyl as R<sub>7</sub> in formula (1) is halogen, hydroxyl or amino.